Cancel claims 1-15, 18, 21, 22, 24-32, and 34-43 and add the following claims 44 through 69.

- 44. The method of Claim 19, wherein Val⁸-GLP-1(7-37)OM is delivered to lower airways of the patient.
- 45. The method of Claim 44, wherein Val⁸-GLP-1/7-37)OH is deposited in the alveoli.
- 46. The method of Claim 19, wherein Val⁸-GLP-1(7-37)OH is inhaled through the mouth of the patient.
- 47. The method of Claim 19, wherein Val⁸-GLP-1(7-37)OH is administered as a pharmaceutical formulation comprising the Val⁸-GLP-1(7-37)OH in a pharmaceutically acceptable carrier.
- 48. The method of Claim 47, wherein the formulation is selected from the group consisting of a solution in an aqueous medium and a suspension in a non-aqueous medium.
- 49. The method of Claim 48, wherein the formulation is administered as an aerosol.
- 50. The method of Claim 47, wherein the formulation is in the form of a dry powder.
- 51. The method of Claim 47, wherein the Val⁸-GLP-1(7-37)OH has a particle size of less than about 10 microns MMAD.
- 52. The method of Claim 51, wherein the Val⁸-GLP-1(7-37)OH has a particle size of about 1 to about 5 microns MMAD.

The method of Claim 52, wherein the Val^8 -GLP-1(7 in the Market 1600 mg/g) 37)OH has a particle size of about 2 to about 3 53. microns MMAD.

- The method of Claim 19, wherein at least about 10% of the Val8-GLP-1(7-37)OH delivered is deposited in 54. the lung.
- The method of Claim 19, wherein the $\text{Mal}^8\text{-GLP-1}(7\text{-}$ 37) OH is delivered from an inhalation device 55. suitable for pulmonary administration and capable of depositing the Val^{8} -GLP-1(7-3V)9H in the lungs of the patient.
 - The method of Claim 55, wherein the device is selected from the group consisting of a nebulizer, a 56. metered-dose inhaler, a dry powder inhaler, and a sprayer.
 - The method of claim 56, wherein the device is a dry 57. powder inhaler.
 - The method of Claim 28, wherein the Val8-GLP-1(7-37)OH is delivered from an inhalation device 58. suitable for pylmonary administration and capable of depositing the Val^8 -GLP-1(7-37)OH in the lungs of the patient.
 - The method of Claim 58, wherein the device is a 59. sprayer for dry powder inhaler.
 - The method of Claim 58, wherein an actuation of the devi¢e administers about 40 μg to about 4,000 μg of 60. Val/8-GLP-1(7-37)OH.
 - The method of Claim 60, wherein an actuation of the 61.

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device administers about 80 μg to about 2,000 μg of Val⁸-GLP-1(7-37)OH.

- 62. The method of Claim 61, wherein an actuation of the device administers about 160 μg to about 1,000 μg of Val⁸-GLP-1(7-37)OH.
- 63. The method of claim 62, wherein an actuation of the device administers about 320 μg to about 500 μg of Val⁸-GLP-1(7-37)OH.
- 64. The method of Claim 33, wherein the Val⁸-GLP-1(7-37)OH is delivered from an inhalation device suitable for pulmonary administration and capable of depositing the Val⁸-GLP-1(7-37)OH in the lungs of the patient.
- 65. The method of Claim 64, wherein the device is a sprayer or dry powder inhaler.
- 66. The method of Claim 64, wherein an actuation of the device administers about 40 μg to about 4,000 μg of Val⁸-GLP-1(7-37)OH.
- 67. The method of Claim 66, wherein an actuation of the device administers about 80 μg to about 2,000 μg of Val⁸-GLP-1(7-37)OH.
- 68. The method of Claim 67, wherein an actuation of the device administers about 160 μg to about 1,000 μg of Val⁸/GLP-1(7-37)OH.
- 69. The method of claim 68, wherein an actuation of the device administers about 320 μg to about 500 μg of Val⁸-GLP-1(7-37)OH.

Py A